

**AMENDEMENTS TO THE CLAIMS**

Claims 1-32 (Canceled)

33. (Previously presented) A transgenic mouse whose genome comprises a homozygous disruption in endogenous mouse brain-specific membrane anchored protein (BSMAP) gene, wherein the transgenic mouse exhibits increased prepulse inhibition, relative to a wild-type mouse.

34. (Previously presented) The transgenic mouse of claim 33, wherein the increased prepulse inhibition is observed after a 100 decibel prepulse.

35. (Previously presented) A cell or tissue obtained from the transgenic mouse of claim 33.

36. (Canceled)

37. (Canceled)

38. (Previously presented) A method of producing a transgenic mouse comprising a disruption in endogenous mouse BSMAP gene, the method comprising:

- (a) introducing a targeting construct capable of disrupting mouse BSMAP gene into a mouse embryonic stem cell;
  - (b) introducing the resulting mouse embryonic stem cell into a blastocyst;
  - (c) implanting the resulting blastocyst into a pseudopregnant mouse, wherein the pseudopregnant mouse gives birth to a chimeric mouse; and
  - (d) breeding the chimeric mouse to produce the transgenic mouse;
- wherein where the disruption is homozygous, the transgenic mouse exhibits increased prepulse inhibition, relative to a wild-type mouse.

39. (Previously presented) The transgenic mouse produced by the method of claim 38.

40. (Previously presented) A targeting construct capable of disrupting a mouse BSMAP gene, the targeting construct comprising:

- (a) a first polynucleotide sequence homologous to a mouse BSMAP gene;
- (b) a second polynucleotide sequence homologous to the mouse BSMAP gene; and
- (c) a selectable marker.

wherein the targeting construct, when introduced into a mouse embryonic stem cell, results in production of a transgenic mouse whose genome comprises a disruption in the mouse BSMAP gene, wherein the transgenic mouse exhibits increased prepulse inhibition, relative to a wild-type mouse.

41. (Previously presented) The targeting construct of claim 40, wherein the targeting construct further comprises a screening marker.
42. (Previously presented) A method of producing a targeting construct capable of disrupting a mouse BSMAP gene, the method comprising:
  - (a) providing a first polynucleotide sequence homologous to a mouse BSMAP gene;
  - (b) providing a second polynucleotide sequence homologous to the mouse BSMAP gene;
  - (c) providing a selectable marker; and
  - (d) inserting the first sequence, second sequence and selectable marker into a vector to produce the targeting construct,  
wherein the targeting construct, when introduced into a mouse embryonic stem cell, results in production of a transgenic mouse whose genome comprises a disruption in the mouse BSMAP gene, wherein the transgenic mouse exhibits increased prepulse inhibition, relative to a wild-type mouse.
43. (Previously presented) A method of producing a targeting construct capable of disrupting a mouse BSMAP gene, the method comprising:
  - (a) providing a polynucleotide comprising a first sequence homologous to a first region of a mouse BSMAP gene and a second sequence homologous to a second region of the mouse BSMAP gene; and
  - (b) inserting a positive selection marker between the first and second sequences to form the targeting construct,  
wherein the targeting construct, when introduced into a mouse embryonic stem cell, results in production of a transgenic mouse whose genome comprises a disruption in the mouse BSMAP gene, wherein the transgenic mouse exhibits increased prepulse inhibition, relative to a wild-type mouse.
44. (Currently amended) A murine embryonic stem cell transformed with the targeting construct of claim 40, wherein the embryonic stem cell produces a disruption in the BSMAP gene in a transgenic mouse which results in a phenotype of increased prepulse inhibition, relative to a wild-type mouse.
45. (Previously presented) A method of identifying an agent that modulates prepulse inhibition, the method comprising:

- (a) administering a test agent to a transgenic mouse comprising a homozygous disruption in endogenous mouse BSMAP gene, wherein the transgenic mouse exhibits increased prepulse inhibition; and
- (b) determining whether the test agent modulates prepulse inhibition in the transgenic mouse.

46. (Currently amended) A method of identifying a potential therapeutic agent for the treatment of schizophrenia, the method comprising:

- (a) administering the potential therapeutic agent to a transgenic mouse comprising a homozygous disruption in endogenous mouse BSMAP gene, wherein the transgenic mouse exhibits increased prepulse inhibition; and
- (b) determining whether the potential therapeutic agent modulates prepulse inhibition in the transgenic mouse, wherein modulation of ~~seizure susceptibility-prepulse inhibition~~ identifies a potential therapeutic agent for the treatment of schizophrenia.